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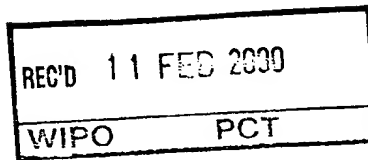
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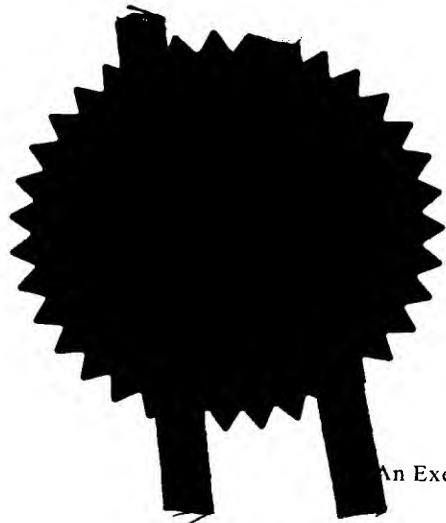
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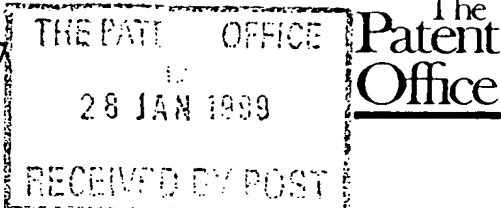
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Cardiff Road
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1. Your reference

IPD/P2352

2. Patent application number

9901764.2

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28 JAN 1999

3. Full name, address and postcode of the or of each applicant (underline all surnames)

The Secretary of State for Defence
Defence Evaluation and Research Agency
Ively Road
Farnborough, Hampshire, GU14 0LX

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

GB

54510008

4. Title of the invention

FIRE DETECTION METHOD

5. Name of your agent (if you have one)

A O Bowdery

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

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Formalities Section
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Country

priority application number
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Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number or earlier application

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8. Is a statement of inventorship and of right if to grant of a patent required in support of this request? (Answer 'Yes' if:

yes

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an applicant, or
- c) any named applicant is a corporate body. See note (d))

Patents Form 1/77

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Continuation sheets of this form

Description 6

Claim(s) 2

Abstract 1

Drawing(s) 3 + 3

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Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77) 1

Request for substantive examination (Patents Form 10/77)

Any other documents (please specify)

11. I / We request the grant of a patent on the basis of this application.

S R Skelton

Signature

SR Skelton

Date 27/01/99

12. Name and daytime telephone number of person to contact in the United Kingdom Karen Lo Sciuto 0117 9132863

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Fire Detection Method

The present invention relates to a method for the detection of overheating of materials in particular electrical equipment, which precedes the onset of fire. Thus the invention is useful in providing advance warning of conditions likely to lead to fire so that preventative action can be taken.

Early warning fire detectors often rely on the detection of smoke particles to trigger an alarm. At this stage, fire is imminent if not already underway and so it is generally too late for preventative action.

Methods for the early detection of fire based upon the vapour detection have also been described. These have potential to provide advanced warning of an imminent fire. These chemical sensing techniques are often based on chemical coatings which interact with the outgassing vapours either through a chemical reaction (USP 5065140) or adsorption. In the former case, the detector lacks versatility in that it reacts only to the vapour of interest. The latter method lacks specificity and requires considerable signal processing effort.

Ion mobility spectrometers are well known in the detection of chemical warfare agents, explosives, propellants, and industrial pollutants. The principles by which they operate and the design of spectrometers are described for example by W. Carr (Ed). "Plasma Chromatography", Plenum Press, London, 1984, and Turner et al., Trends in Analytical Chemistry, 13, 7 (1994) 275-280. In essence, an ion mobility spectrometer consists of an ionisation region coupled to an ion drift tube via a shutter grid. A sample is introduced into the ionisation region together with a carrier gas (such as air), for example using a suction pump. In the ionisation region, the carrier gas molecules (as well as any sample) are ionised by beta radiation from a Ni-63 radioactive source, or other

methods such as corona discharge or photoionisation. Reactant ions are produced from the carrier gas (such as air) which react with the sample gas, generally in a complex manner, so as to result in product ions. Under the influence of an applied electric field, reactant and product ions are extracted from the ionisation region into the ion drift region. In the ion drift region, the ions separate due to their different mobilities determined by their size, charge and polarisability. They are collected at a collector electrode where they are neutralised, and so generate an electric current that can be measured. Data is generated at a rapid rate. Repeat scans are suitably averaged to improve the signal to noise ratio.

The spectrometer can be arranged to detect either positive or negative ions by reversal of the voltages.

The present invention provides a method for detecting gases or vapours emitted from materials under conditions at which there is a risk of the onset of fire, said method comprising sampling gas from the region of the material using an ion mobility spectrometer, detecting the ion peaks of volatilised material.

The advantages of the use of ion mobility spectrometry in this application are associated with its extreme sensitivity and selectivity. The spectrometer can be pre-set, by controlling the potentials applied to the drift region for example, so as to detect selected target gases which are emitted during outgassing through heating in any specific environment. The selectivity of the device means that it could be applied to a variety of different environments. However, it will be particularly useful in high technology environments, such as data processing and computer facilities, telephone exchanges, space stations, industrial plants especially chemical plants or plants which deal with inflammable materials, where the

risk of fire as a result of overheating devices and in particular electrical components is high, and the consequences of such a fire are extreme in economic terms at least.

- 5 Sampling can take place either continuously or at intervals, for example at pre-set intervals. Preferably, the sampling will be carried out continuously, as this allows an increasing signal, produced as a result of increased outgassing of a particular gas from the materials, for example as the
10 temperature rises, to be detected against background noise levels.

- The spectrometer may be connected to a warning or alarm system that may be triggered automatically, using various
15 predetermined parameters. For example, if a signal peak reaches a particular intensity, this may trigger the alarm. An alternative, particularly suitable with continuous monitoring, would be to arrange for the alarm to be triggered when any signal peak increases significantly over a period of
20 time. This may be indicative of a rise in temperature of the component that gives rise to the particular gas or vapour.

- The alarm may be connected to the detection of features characteristic of thermal degradation and not just simple
25 over-heating of components. This would ensure that only potentially serious situations resulted in an actual alarm, reducing false positives to a minimum.

- The levels and the parameters used to trigger the alarm will
30 vary depending upon the particular circumstances in which the spectrometer is being employed. Again, these can be determined using routine methods and the control systems designed appropriately.

- 35 There is no need to know or analyse the materials which are emitted, provided it can be ensured that at least some of

- these fall within the detection range set on the ion mobility spectrometer. This can be done by routine methods. For example, one or more representative components present in the particular environment to be monitored can be heated under
- 5 safe test conditions and the signal generated as a result of the emission of material monitored. The detection range of the ion mobility spectrometer can then be adjusted to ensure a signal is generated under these conditions.
- 10 The spectrometer will be pre-set to detect either positive or negative ions depending upon the nature of the signal in each case. Selection of the most sensitive signal in each particular case can be determined, again by routine methods.
- 15 Many solid materials that release volatile material when heated, but in particular are electrical components such as printed circuit boards, resistors and lacquer-coated materials. The spectrometer will suitably be set to detect vapours emitted from such components.
- 20 Available ion mobility spectrometers are convenient to use in that it is small and hand-held. For use in the context of the invention however, size and portability may be less critical. The spectrometer may be installed in the
- 25 environment on a permanent basis. It would not, under these circumstances be required to be as robust as a device which is intended, for example to be carried onto a battle-field. This may result in cost savings in the spectrometer itself.
- 30 The invention will now be particularly described by way of example with reference to the accompanying diagrammatic drawings in which:

Figure 1 is a schematic diagram of an ion mobility

35 spectrometer;

Figure 2 shows the positive ion mobility spectra for a heated lacquer-coated printed circuit board at temperatures of from 50 to 85°C;

- 5 Figure 3 shows the positive ion mobility spectra for a heated lacquer-coated printed circuit board at temperatures of from 85 to 105°C;

- Figure 4 shows the negative ion mobility spectra for a
10 resistor heated to 50 to 100°C; and

Figure 5 shows the positive ion mobility spectra for a resistor heated to 50 to 100°C.

- 15 The illustrated device in Figure 1 comprises an inlet system comprising a heated nozzle (1) and a silicon rubber membrane (2). Gas sample is admitted through the inlet system as a result of the action of a diaphragm pump (3) operated by a motor (4). Sample transfers into an ionisation section (5)
20 where a nickel-63 ion source generates the ions. A pulse of ions (generally about 0.2ms) is admitted into a drift tube section (6) by manipulation of the potentials on a grid assembly (7). The drift tube (6) is typically about 4cm long with an electric field of 250V/cm. Ions pass to a collector
25 electrode (8), where they are neutralised, generating a current which is passed to a microprocessor (9), which generates a signal, if necessary after amplification. The signal may be passed to a display assembly (10).
- 30 In general ion drift times are less than 20ms, and so spectra may be gathered at the rate of say 50Hz.

Example 1

- A lacquer coated printed circuit board was heated from 50 to
35 105°C in the vicinity of an ion mobility spectrometer that was operational throughout. At the start of the heating process,

the positive reactant ion peak (i.e. that produced as a result of the ionisation of air), is the major feature of the spectrum (see Figure 2 - trace at the back of the representation). As the printed circuit board was heated, 5 this peak is replaced by an ion of reduced mobility (further to the right in the representation), formed by the reaction of vapour emitted by the board with reactant ions in the instrument. This characteristic feature increases in intensity and then falls as a further prominent ion is formed 10 (Figure 3).

Example 2

Example 1 was repeated but this time with a resistor in place of the printed circuit board. In a first experiment, the 15 resistor was heated from 50 to 100°C and in a second experiment, a temperature range of from 90 to 140°C was used. Significant changes in the negative ion spectra (Figures 4 and 5 respectively) were recorded.

20 The sensitivity of this technique is clear from this example, as resistors do not generally emit volatile materials.

Claims

1. A method for detecting gases or vapours emitted from materials under conditions at which there is a risk of the onset of fire, said method comprising sampling gas from the region of the material using an ion mobility spectrometer, detecting the ion peak of volatilised material.
2. A method according to claim 1 which is effected in a data processing or computer facility, a telephone exchange, a space station, or an industrial plant.
3. A method according to claim 1 or claim 2 wherein sampling is effected continuously.
4. A method according to claim 1 or claim 2 wherein sampling is effected at predetermined time intervals.
5. A method according to any one of the preceding claims wherein the ion mobility spectrometer is connected to an alarm system.
6. A method according to claim 5 wherein the alarm is triggered when an ion peak reaches a predetermined intensity level.
7. A method according to claim 5 wherein the alarm is triggered when any ion peak increases significantly over a period of time.
8. A method according to any one of claims 5 to 7 wherein the alarm is triggered when a feature characteristic of thermal degradation is detected.

9. A method according to any one of the preceding claims wherein the ion mobility spectrometer is set to detect volatile material released from an electrical component.

5 10. A method according to claim 9 wherein the electrical component is a printed circuit board or a resistor.

11. The use of an ion mobility spectrometer in the detection of gases or vapours emitted from materials under conditions at
10 which there is a risk of the onset of fire.

Abstract

A method for detecting gases or vapours emitted from materials, such as electrical components, under conditions at which there is a risk of the onset of fire, said method comprising sampling gas from the region of the material using an ion mobility spectrometer tuned to detect specific volatilised materials. The signal can be used to determine a fire risk and to trigger alarms as necessary. The method provides rapid detection with high sensitivity and high selectivity so as to minimise false alarms.

FIGURE 1

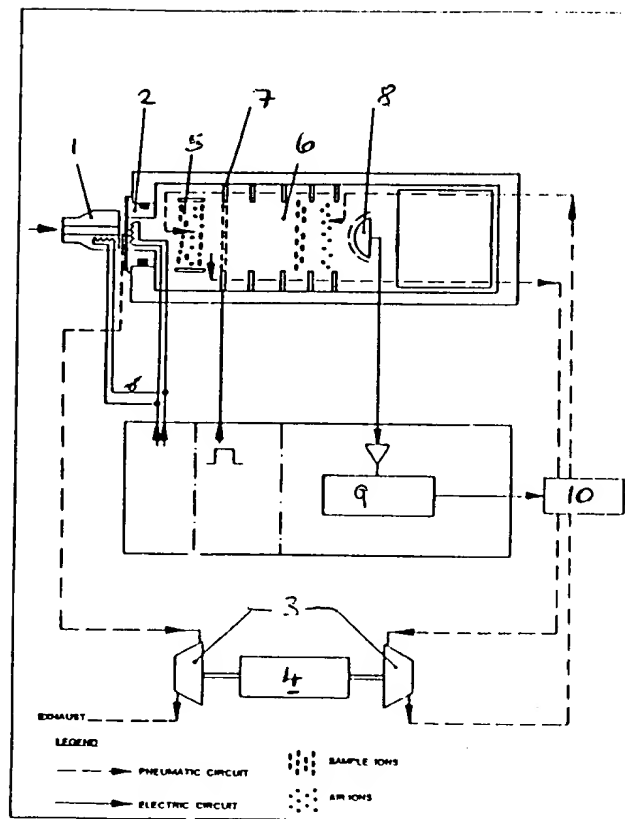


FIGURE 2

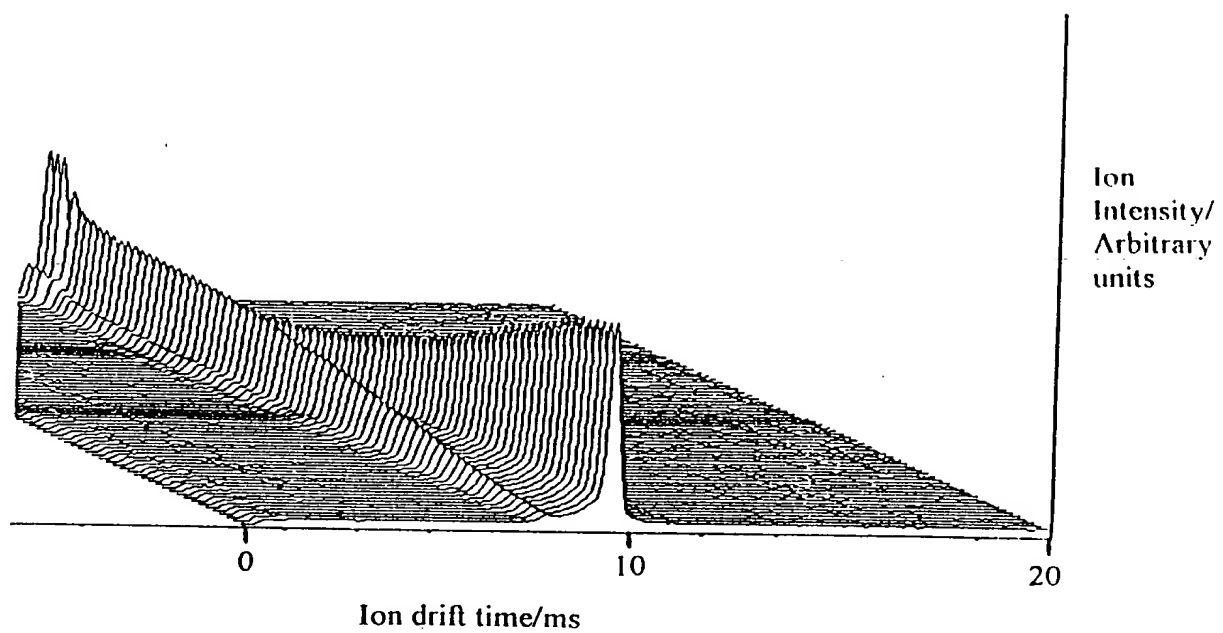


FIGURE 3

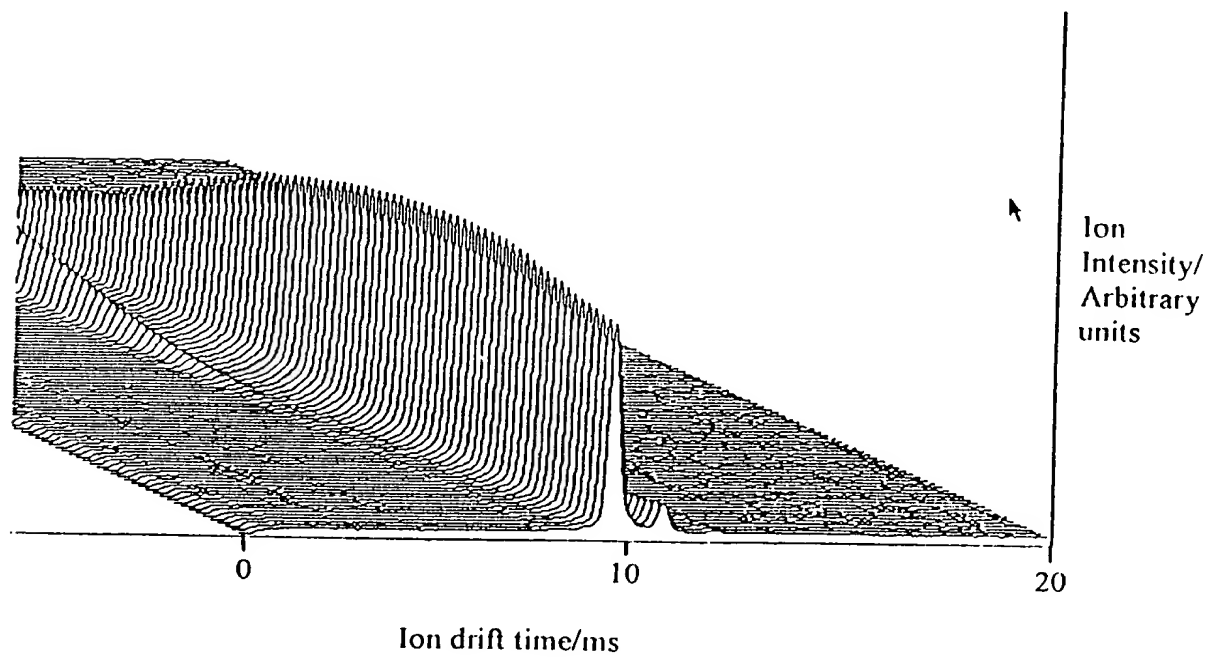


FIGURE 4

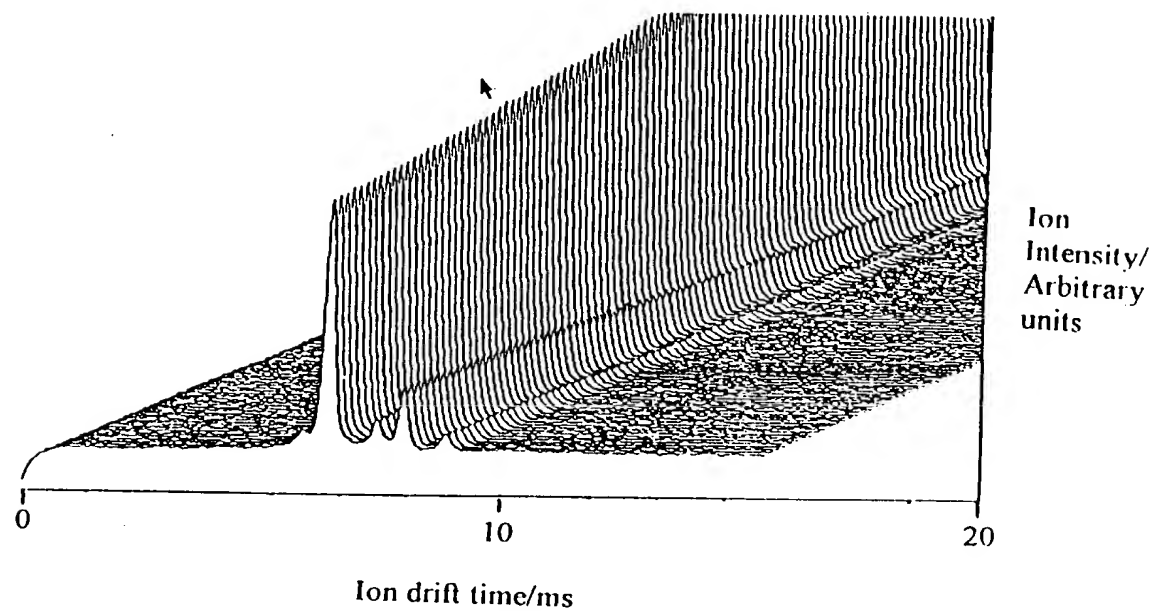
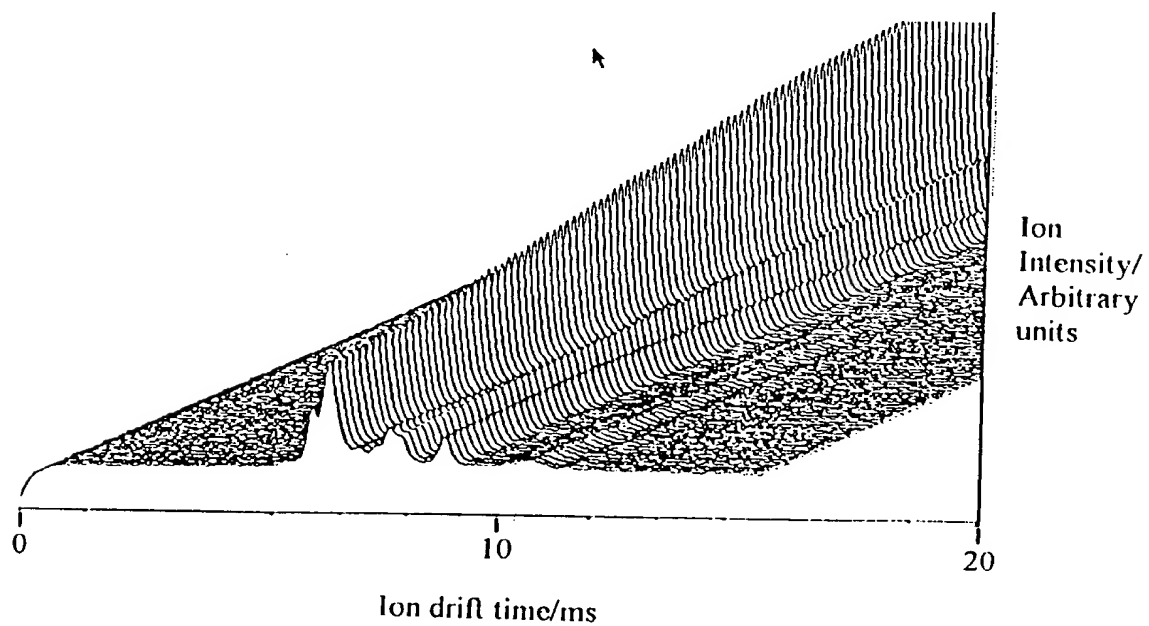


FIGURE 5



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